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that it has a cytolytic activity on pathogenic cells, said pathogenic cells being cells which are non-naturally occurring within the body consisting of microbial pathogenic organisms and malignant cells; and it is non-hemolytic, namely it has no cytolytic effect on red blood cells or has a cytolytic effect on red blood cells at concentrations which are substantially higher than that in which it manifests said cytolytic activity on pathogenic cells, said non-hemolytic cytolytic peptide being selected from the group consisting of:

- (A) a cyclic derivative of a peptide having a net positive charge which is greater than +1, and comprising both L-amino acid residues and D-amino acid residues, or comprising only D-amino acid residues, and comprising an α -helix breaker moiety;
- (B) a peptide comprising both L-amino acid residues and D-amino acid residues, having a net positive charge which is greater than +1, and having a sequence of amino acids such that the same amino acid sequence in which each residue is in the L-configuration is not found in nature, and cyclic derivatives thereof; and
- (C) a random copolymer consisting of a hydrophobic, a positively charged and a D-amino acid, with the proviso that the peptide is not that of SEQ ID NO:1.

14 (Five-Times-Amended). A cyclic derivative of a non-natural synthetic peptide according to claim 7, selected from the group of peptides consisting of those herein designated 92-95 (SEQ ID NOS:92-95, respectively), of the sequence:

92) Cyclic Cys Lys Leu Leu Leu Lys Leu Leu Leu Lys Leu Leu Lys Cys,

93) Cyclic Cys Lys Leu Leu Leu Lys Leu Lys Leu Lys Leu Lys Cys,

94) HN - Lys Leu Leu Leu Lys Leu Leu Leu Lys Leu Leu Lys - CO, and

95) HN - Lys Leu Leu Leu Lys Leu Lys Leu Lys Leu Leu Lys - CO.

31 (Twice-Amended). A composition comprising a pharmaceutically acceptable carrier and a peptide according to claim 1 in an anti-viral effective amount.

32 (Amended). The composition of claim 31, wherein said antiviral effective amount is an amount effective to inhibit viral-induced hemolysis.

REMARKS

Claims 1-14, 20, 21, 27-35 and 37 presently appear in this case. Claims 1-14, 20, 21 and 27-34 have been rejected. Claims 35 and 37 have been objected to and, presumably, would be allowable if rewritten into independent